

I. Amendments to the Claims:

This Listing of the Claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

Claim 1. (currently amended) A pharmaceutical composition comprising about 5 to about 20 mg hydrocodone or pharmaceutically acceptable salt thereof and 0.055 to 0.56 mg naltrexone or pharmaceutically acceptable salt thereof, said naltrexone or pharmaceutically acceptable salt thereof and said hydrocodone or pharmaceutically acceptable salt thereof in a ratio of from 0.011:1 to 0.0125:1 0.028:1.

Claim 2. (currently amended) The pharmaceutical composition of claim 1 comprising about 5 mg hydrocodone or pharmaceutically acceptable salt thereof and ~~from 0.055 mg to 0.14 mg~~ of naltrexone or pharmaceutically acceptable salt thereof.

Claim 3. (currently amended) The pharmaceutical composition of claim 1 comprising about 7.5 mg hydrocodone or pharmaceutically acceptable salt thereof and ~~from 0.0825 mg to 0.21 mg~~ of naltrexone or pharmaceutically acceptable salt thereof.

Claim 4. (currently amended) The pharmaceutical composition of claim 1 comprising about 10 mg hydrocodone or pharmaceutically acceptable salt thereof and ~~from 0.11 mg to 0.28 mg~~ of naltrexone or pharmaceutically acceptable salt thereof.

Claim 5. (currently amended) The pharmaceutical composition of claim 1 comprising about 15 mg hydrocodone or pharmaceutically acceptable salt thereof and ~~from 0.165 mg to 0.42 mg~~ of naltrexone or pharmaceutically acceptable salt thereof.

Claim 6. (currently amended) The pharmaceutical composition of claim 1 comprising about 20 mg hydrocodone or pharmaceutically acceptable salt thereof and ~~from 0.22 mg to 0.56 mg~~ of naltrexone or pharmaceutically acceptable salt thereof.

Claim 7. (original) The pharmaceutical composition of claim 1 comprising about 5 mg hydrocodone or pharmaceutically acceptable salt thereof and 0.0625 mg of naltrexone or pharmaceutically acceptable salt thereof.

Claim 8. (original) The pharmaceutical composition of claim 1 comprising about 7.5 mg hydrocodone or pharmaceutically acceptable salt thereof and 0.09375 mg of naltrexone or pharmaceutically acceptable salt thereof.

Claim 9. (original) The pharmaceutical composition of claim 1 comprising about 10 mg hydrocodone or pharmaceutically acceptable salt thereof and 0.125 mg of naltrexone or pharmaceutically acceptable salt thereof.

Claim 10. (original) The pharmaceutical composition of claim 1 comprising about 15 mg hydrocodone or pharmaceutically acceptable salt thereof and 0.1875 mg of naltrexone or pharmaceutically acceptable salt thereof.

Claim 11. (original) The pharmaceutical composition of claim 1 comprising about 20 mg hydrocodone or pharmaceutically acceptable salt thereof and 0.25 mg of naltrexone or pharmaceutically acceptable salt thereof.

Claim 12. (original) The pharmaceutical composition of claim 1 further comprising a sustained release excipient which provides a sustained release of the hydrocodone or pharmaceutically acceptable salt thereof.

Claim 13. (original) The pharmaceutical composition of claim 1 further comprising a sustained release excipient which provides a sustained release of the naltrexone or pharmaceutically acceptable salt thereof.

Claim 14. (original) The pharmaceutical composition of claim 1 further comprising a sustained release excipient which provides a sustained release of the hydrocodone or pharmaceutically acceptable salt thereof and the naltrexone or pharmaceutically acceptable salt thereof.

Claim 15. (currently amended) The pharmaceutical composition of claim 12, wherein the ~~dosage form composition~~ provides effective pain relief for at least 12 hours after steady state oral administration to human patients.

Claim 16. (currently amended) The pharmaceutical composition of claim 12, wherein the ~~dosage form composition~~ provides effective pain relief for at least 24 hours after steady state oral administration to human patients.

Claim 17. (currently amended) The pharmaceutical composition of claim 14, wherein the hydrocodone or pharmaceutically acceptable salt thereof and the naltrexone or pharmaceutically acceptable salt thereof are substantially interdispersed in said sustained release excipient.

Claim 18. (currently amended) The pharmaceutical composition of claim 1, wherein said hydrocodone is in the form of the bitartrate salt.

Claim 19. (currently amended) The pharmaceutical composition of claim 1, wherein said naltrexone is in the form of the hydrochloride salt.

Claim 20. (previously presented) The pharmaceutical composition of claim 1 further comprising a non-steroidal anti-inflammatory drug selected from the group consisting of ibuprofen, diclofenac, naproxen, benoxaprofen, flurbiprofen, fenoprofen, flubufen, ketoprofen, indoprofen, piroprofen, carprofen, oxaprozin, pramoprofen, muroprofen, trioxaprofen, suprofen, aminoprofen, tiaprofenic acid, fluprofen, bucloxic acid, indomethacin, sulindac, tolmetin, zomepirac, tiopinac, zidometacin, acemetacin, fentiazac, clidanac, oxpinac, mefenamic acid, meclofenamic acid, flufenamic acid, niflumic acid, tolfenamic acid, diflurisal, flufenisal, piroxicam, sudoxicam, isoxicam, pharmaceutically acceptable salts thereof and mixtures thereof

Claim 21. (previously presented) A method of treating pain in a human patient comprising orally administering a pharmaceutical composition according to claim 1.

Claim 22. (currently amended) A method of preparing a pharmaceutical composition comprising combining about 5 to about 20 mg hydrocodone or pharmaceutically acceptable salt thereof and 0.055 to 0.56 mg naltrexone or pharmaceutically acceptable salt thereof ~~into~~ into an oral dosage form, said naltrexone or pharmaceutically acceptable salt thereof and said hydrocodone or pharmaceutically acceptable salt thereof in a ratio of from 0.011:1 to 0.0125:1 0.028:1.

Claim 23. (previously presented) A method of deterring abuse of a hydrocodone formulation comprising preparing a pharmaceutical formulation of claim 1.

Claim 24. (currently amended) A method of treating pain in a human patient comprising administering a dosage form of claim 1 to said patient The use of hydrocodone or a pharmaceutically acceptable salt thereof, in the preparation of a dosage form according to claim 1.

Claim 25. (currently amended) The method of claim 24, wherein the dosage form is administered once-a-day use of naltrexone or a pharmaceutically acceptable salt thereof, in the preparation of a dosage form according to claim 1.

Claim 26. (currently amended) The method of claim 24, wherein the dosage form is administered twice-a-day use of hydrocodone or a pharmaceutically acceptable salt thereof, and naltrexone or a pharmaceutically acceptable salt thereof, in the preparation of a dosage form according to claims 1.

Claim 27 (new): A pharmaceutical composition comprising hydrocodone or pharmaceutically acceptable salt thereof and naltrexone or pharmaceutically acceptable salt thereof, wherein the

naltrexone or pharmaceutically acceptable salt thereof and said hydrocodone or pharmaceutically acceptable salt thereof are in a ratio of 0.0125:1.

Claim 28 (new): The pharmaceutical composition of claim 27, wherein the drugs in the composition consist of said naltrexone or pharmaceutically acceptable salt thereof and said hydrocodone or pharmaceutically acceptable salt thereof.

Claim 29 (new): The pharmaceutical composition of claim 27 comprising from about 5 to about 20 mg of said hydrocodone or pharmaceutically acceptable salt thereof.